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Sterling Medivations, Inc.
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Los Altos Hills, CA 94022
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510(k) SUMMARY

Date Submitted: March 19, 2001

Submitter: Sterling Medivations, Inc. 25285 La Loma Drive, Los Altos Hills, CA 94022
Company Phone 650-949-0470, Company Fax 650-949-0342

Contact: Joel Douglas, Chief Technology Officer
Sterling Medivations, Inc.
Applicant Phone 650-817-4083 or 650-949-0470
Applicant Fax 408-297-9473 or 650-949-0342

Trade Name of Device: Simplicity™ Easy Soft Infusion Set for infusion and/or injection of fluids into the body below the surface of the skin when attached to an external pump or syringe.

Common Name of Device: Intravascular administration set.
Classification Name: Percutaneous intravascular catheter.

Predicate Device: Simplicity Easy Access Infusion Set Infusion Set, FDA 510 (k) K014062 .

Description of the New Device: Sterling Medivations Inc.'s ("SMI") Simplicity Easy Soft Infusion Set is designed for infusion and/or injection of fluids into the body below the surface of the skin when attached to an external pump or syringe.

The Simplicity Easy Soft Infusion Set proposed for commercial distribution is similar in all significant respects to the existing Simplicity Easy Access Infusion Set Infusion Set, FDA 510 (k) K014062 and it has the same intended use.

The device consists of four main parts: (1) infusion catheters are made from Polytetrafluoroethylene (PTFE), (2) an infusion hub that provides the patient the capability of disconnecting the connecting tube from the infusion catheter, (3) a connecting tube and threaded reservoir connector and (4) an insertion needle ring.

The Simplicity Easy Soft Infusion Set is an infusion administration set, connecting to a medicine reservoir or syringe that is placed in an external infusion device and inserted below the surface of the skin of a patient. The SMI Simplicity Easy Soft Infusion Set may be used with any infusion device that delivers continuous or intermittent flow.

The administration set attaches to the reservoir/syringe by means of a threaded luer connector, and under the surface of the skin in the patient through an indwelling cannula formed from PTFE,. The connecting tubing is made from a polyethylene tube.

The PTFE indwelling cannula are inserted with insertion needles formed from AISI 304 stainless steel are introduced below the surface of the skin. The insertion needle is removed and the connector needle is attached to the hub fixed to the indwelling catheter. This seal on the connector needle mates with the indwelling catheter hub that forms a seal that permits the infusion of medication without leakage. The connector needle is made from AISI 304 stainless steel and it is connected to the connecting tubing. The connector tubing proximal end is attached to a threaded luer connector for attachment to the medicine reservoir.

Intended Use of the New Device: The intended use of the Simplicity Easy Soft Infusion Set is to provide a means to for infusion and/or injection of fluids into the body below the surface of the skin when attached to an external pump or syringe.

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Comparison of the Technological Features of the New Device and Predicate Device:

The Simplicity Easy Soft Infusion Set proposed for commercial distribution is similar in all significant respects to the existing Simplicity Easy Access Infusion Set Infusion Set, FDA 510 (k) K014062 and it has the same intended use.

The materials and manufacturing processes are substantially equivalent, the labeling is substantially equivalent and it has the same intended use as the Simplicity Easy Access Infusion Set Infusion Set, FDA 510 (k) K014062.

The differences that exist between the new and predicate device are as follows:

1. The new device has 360 degree rotating infusion hub..

2. The new device has multiple cannula form from polytetrafluorthylene (PTFE) and the predicate device has has multiple indwelling catheters formed from AISI 304 stainless steel

Performance Data Supporting Substantial Equivalence: To prove substantial equivalence the Simplicity Easy Soft Infusion Set meets the catheter requirements of:

- CDRH 21 C.F.R. Section 880.5440 Intravascular administration set
- ISO 10555 Sterile, single use intravascular catheters (Part 1: General Requirements)
- ISO 10555 Sterile, single use intravascular catheters (Part 5: Peripheral Catheters).
- ISO 11135:1994 Medical devices -- Validation and routine control of ethylene oxide sterilization
- ISO 11138-2:1994 Sterilization of health care products -- Biological indicators -- Part 2: Biological indicators for ethylene oxide sterilization.
- ISO 594-1:1986 Conical fittings with a 6 % (Luer) taper for syringes, needles and certain other medical equipment -- Part 1: General requirements,
- ISO 594-2:1998 Conical fittings with 6 % (Luer) taper for syringes, needles and certain other medical equipment -- Part 2: Lock fittings
- ISO 9626: 1991 Stainless Steel needle tubing for the manufacture of medical devices.
- ISO 11607: 1997 Packaging for terminally sterilized medical devices.
- ISO 8535: 1991 Sterile single use syringes, with or without needle, for insulin.
- FDA Guidelines on validation of the Limulus Amebocyte Lysate (LAL) Test as an end product endotoxin test for human and animal parenteral drugs, biological products, and medical devices.
- ODE Blue Book Memorandum #K90-1.
- ISO 10993-1, Biological evaluation of medical devices - Part 1: Evaluation and testing.

And the design process adhered to is the Center for Devices and Radiological Health. DESIGN CONTROL GUIDANCE FOR MEDICAL DEVICE MANUFACTURERS. This Guidance relates to FDA 21 CFR 820.30 and Sub-clause 4.4 of ISO 9001. This is substantially equivalent to the predicate device.

Signed,

Joel S. Douglas
Chief Technology Officer

DEPARTMENT OF HEALTH & HUMAN SERVICES

Public Health Service

Food and Drug Administration
9200 Corporate Boulevard
Rockville MD 20850

APR 1 0 2002

Mr. Joel Douglas
Chief Technology Officer
Sterling Medivations, Incorporated
25285 La Loma Drive
Los Altos Hills, California 94022-4583

Re: K020912
Trade/Device Name: Simplicity Easy Soft Infusion Set
Regulation Number: 880.5440
Regulation Name: Intravascular Administration Set
Regulatory Class: II
Product Code: FPA
Dated: March 19, 2002
Received: March 20, 2002

Dear Mr. Douglas:

We have reviewed your Section 510(k) premarket notification of intent to market the device referenced above and have determined the device is substantially equivalent (for the indications for use stated in the enclosure) to legally marketed predicate devices marketed in interstate commerce prior to May 28, 1976, the enactment date of the Medical Device Amendments, or to devices that have been reclassified in accordance with the provisions of the Federal Food, Drug, and Cosmetic Act (Act) that do not require approval of a premarket approval application (PMA). You may, therefore, market the device, subject to the general controls provisions of the Act. The general controls provisions of the Act include requirements for annual registration, listing of devices, good manufacturing practice, labeling, and prohibitions against misbranding and adulteration.

If your device is classified (see above) into either class II (Special Controls) or class III (PMA), it may be subject to such additional controls. Existing major regulations affecting your device can be found in the Code of Federal Regulations, Title 21, Parts 800 to 898. In addition, FDA may publish further announcements concerning your device in the Federal Register.

Please be advised that FDA's issuance of a substantial equivalence determination does not mean that FDA has made a determination that your device complies with other requirements of the Act or any Federal statutes and regulations administered by other Federal agencies.

You must comply with all the Act's requirements, including, but not limited to: registration and listing (21 CFR Part 807); labeling (21 CFR Part 801); good manufacturing practice requirements as set forth in the quality systems (QS) regulation (21 CFR Part 820); and if applicable, the electronic product radiation control provisions (Sections 531-542 of the Act); 21 CFR 1000-1050.

This letter will allow you to begin marketing your device as described in your Section 510(k) premarket notification. The FDA finding of substantial equivalence of your device to a legally marketed predicate device results in a classification for your device and thus, permits your device to proceed to the market.

If you desire specific advice for your device on our labeling regulation (21 CFR Part 801 and additionally 21 CFR Part 809.10 for in vitro diagnostic devices), please contact the Office of Compliance at (301) 594-4618 . Additionally, for questions on the promotion and advertising of your device, please contact the Office of Compliance at (301) 594-4639. Also, please note the regulation entitled, "Misbranding by reference to premarket notification" (21CFR Part 807.97). Other general information on your responsibilities under the Act may be obtained from the Division of Small Manufacturers, International and Consumer Assistance at its toll-free number (800) 638-2041 or (301) 443-6597 or at its Internet address http://www.fda.gov/cdrh/dsma/dsmamain.html

Sincerely yours,

Timothy A. Ulatowski
Director
Division of Dental, Infection Control
and General Hospital Devices
Office of Device Evaluation
Center for Devices and
Radiological Health

Enclosure

510(k) Number (if known):

Device Name: Simplicity Easy Soft Infusion Set

Indications For Use:

The intended use of the Simplicity Easy Soft Infusion Set is to provide a means for infusion and/or injection of fluids into the body below the surface of the skin when attached to an external pump or syringe.

(PLEASE DO NOT WRITE BELOW THIS LINE-CONTINUE ON ANOTHER PAGE IF NEEDED)

Concurrence of CDRH, Office of Device Evaluation (ODE)

Prescription Use ✓ OR Over-The-Counter Use ___
(PER 21 CFR 801.109)

(Optional Format 1-2-96)

Patricia Cascante
(Division Sign-Off)
Division of Dental, Infection Control,
and General Hospital Devices
510(k) Number K020912